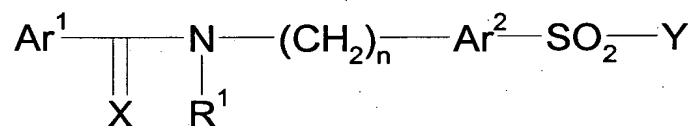


Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) Sulfonamide compounds according to formula I



I

with its geometrical isomers, in an optically active form as enantiomers, diastereomers, as well as in the form of racemates, as well as pharmaceutically acceptable salts thereof, wherein

Ar¹ and Ar² are independently from each other substituted or unsubstituted aryl or heteroaryl groups,

X is O or S;

R¹ is hydrogen or a C₁-C₆-alkyl ~~or hydroxy~~ group, or R¹ forms a substituted or unsubstituted 5-6-membered saturated or unsaturated ring with Ar¹;

n is an integer from 0 to 5;

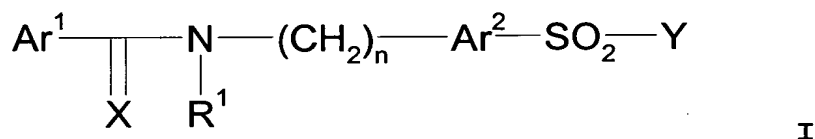
Y within formula I is a substituted or unsubstituted piperidino moiety, whereby one nitrogen atom within ~~said piperidine~~ said piperidino moiety forms a bond with the sulfonyl group of formula I thus providing a sulfonamide,

with the proviso that if Ar¹ is 4-chlorophenyl, X is O, R¹ is H, Ar² is thienyl, and Y is not para-substituted by 2-hydroxyethyl,

with the further proviso that if Ar¹ is phenyl, X is O, R¹ is H, Ar² is thienyl, and Y is not substituted in its β

position by a benzo [5,6] cyclohepta [1,2b] pyridine or a benzo [5,6] cyclohept [3,4]ene [1,2b] pyridine with the final proviso that if X is oxygen and Y is a 4-8 membered saturated cyclic alkyl containing one or two nitrogen atoms, Y shall not be substituted by a group (C=O)N(R,R') at the α -position of the sulfonamide nitrogen.

2. (Currently Amended) A composition for treatment of disorders associated with the abnormal expression or activity of JNK comprising a sulfonamide compound according to formula I



with its geometrical isomers, in an optically active form as enantiomers, dia-stereomers, as well as in the form of racemates, as well as pharmaceutically acceptable salts thereof, wherein

Ar¹ and Ar² are independently from each other substituted or unsubstituted aryl or heteroaryl groups,

X is O or S;

R¹ is hydrogen or a C₁-C₆-alkyl group, or R¹ forms a substituted or unsubstituted 5-6-membered saturated or unsaturated ring with Ar¹;

n is an integer from 0 to 5;

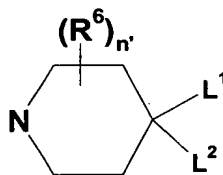
Y within formula I is a piperidino group whereby one nitrogen atom within said piperidino group forms a bond with the sulfonyl group of formula I thus providing a sulfonamide;

with the proviso that if Y is substituted at the β -position of the piperidino- nitrogen by a benzo[5,

6]cyclohepta[1, 2b]pyridine, or a benzo[5, 6]cyclohept-
[3,4]-ene [1, 2b]pyridine, while Ar^2 is thienyl, X is
oxygen, R^1 is hydrogen and n is 1, Ar^1 shall not be a phenyl
group;

and a pharmaceutically acceptable carrier with the
final proviso that if X is oxygen and Y is a 4-8 membered
saturated cyclic alkyl containing one or two nitrogen atoms, Y
shall not be substituted by a group $(\text{C}=\text{O})\text{N}(\text{R}, \text{R}')$ at the α -
position of the sulfonamide nitrogen.

3. (Currently Amended) A sulfonamide compound
according to claim 1 or a composition according to claim 2,
wherein Y is a piperidino group of the formula



whereby, L^1 and L^2 are independently selected from
each other from the group consisting of H, substituted or
unsubstituted C_1 - C_6 -aliphatic alkyl, substituted or
unsubstituted C_2 - C_6 -alkenyl, substituted or unsubstituted C_2 - C_6 -
alkynyl, substituted or unsubstituted cyclic C_4 - C_8 -alkyl
optionally containing 1-3 heteroatoms and optionally fused
with aryl or heteroaryl; or L^1 and L^2 are independently
selected from the group consisting of substituted or
unsubstituted aryl, substituted or unsubstituted heteroaryl,
aryl- C_1 - C_6 -alkyl, heteroaryl- C_1 - C_6 -alkyl, $-\text{C}(\text{O})-\text{OR}^3$, $-\text{C}(\text{O})-\text{R}^3$, $-\text{C}(\text{O})-\text{NR}^3\text{R}^3$, $-\text{NR}^3\text{R}^3$, $-\text{NR}^3\text{C}(\text{O})\text{R}^3$, $-\text{NR}^3\text{C}(\text{O})\text{NR}^3\text{R}^3$, $-(\text{SO})\text{R}^3$, $-(\text{SO}_2)\text{R}^3$, $-\text{NSO}_2\text{R}^3$, $-\text{NHSO}_2\text{R}^3$, and $-\text{SO}_2\text{NR}^3\text{R}^3$,

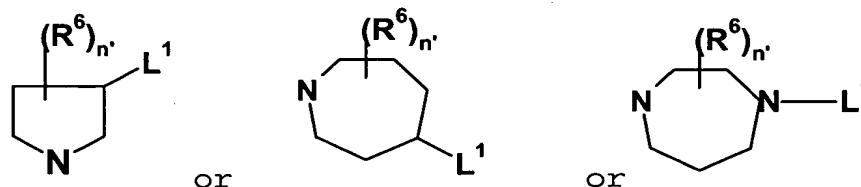
with R^3 and $R^{3'}$ being substituents independently selected from the group consisting of H, substituted or unsubstituted C_1 - C_6 -alkyl, substituted or unsubstituted C_2 - C_6 -alkenyl, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted or unsubstituted aryl- C_1 - C_6 -alkyl and substituted or unsubstituted heteroaryl- C_1 - C_6 -alkyl;

said aryl or heteroaryl groups being optionally substituted C_1 - C_6 -alkyl, C_1 - C_6 -alkoxy, C_2 - C_6 -alkenyl, C_2 - C_6 -alkynyl, amino, acylamino, aminocarbonyl, C_1 - C_6 -alkoxycarbonyl, aryl, carboxyl, cyano, halogen, hydroxy, nitro, sulfonyl, sulfoxy, and C_1 - C_6 -thioalkoxy, and

R^6 is selected from the group consisting of hydrogen, substituted or unsubstituted C_1 - C_6 -alkyl, substituted or unsubstituted C_1 - C_6 -alkoxy, OH, halogen, nitro, cyano, sulfonyl and oxo (=O), and

n' is an integer from 0 to 4.

4. (Withdrawn) A sulfonamide derivative according to claim 1 or 2, wherein Y is a pyrrolidine, an azepan or a 1,4-diazepan moiety of the below formulas



wherein L^1 is selected from the group comprising or consisting of H, substituted or unsubstituted C_1 - C_6 -alkyl, substituted or unsubstituted C_2 - C_6 -alkenyl, substituted or unsubstituted C_2 - C_6 -alkynyl, substituted or unsubstituted cyclic C_4 - C_8 -alkyl optionally containing 1-3 heteroatoms and optionally fused with aryl or heteroaryl; or L^1 and L^2 are

independently selected from the group comprising or consisting of substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, aryl-C₁-C₆-alkyl, heteroaryl-C₁-C₆-alkyl, -C(O)-OR³, -C(O)-R³, -C(O)-NR^{3'}R³, -NR^{3'}R³, -NR^{3'}C(O)R³, -NR^{3'}C(O)NR^{3'}R³, -(SO)R³, -(SO₂)R³, -NSO₂R³, -SO₂NR^{3'}R³;

R³ and R^{3'} are substituents independently selected from the group comprising or consisting of H, substituted or unsubstituted C₁-C₆-alkyl, substituted or unsubstituted C₂-C₆-alkenyl, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted or unsubstituted aryl-C₁-C₆-alkyl, substituted or unsubstituted heteroaryl-C₁-C₆-alkyl;

R⁶ is selected from the group comprising or consisting of hydrogen, substituted or unsubstituted C₁-C₆-alkyl, substituted or unsubstituted C₁-C₆-alkoxy, OH, halogen, nitro, cyano, sulfonyl, oxo (=O), sulfoxy, acyloxy, thioalkoxy and n' is an integer from 0 to 4, preferably 0.

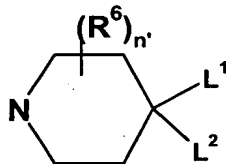
5. (Currently Amended) A sulfonamide compound according to claim 1, wherein Ar¹ and Ar² are independently selected from the group consisting of phenyl, thienyl, furyl, pyridyl, ~~thioxo-dihydropyridine~~ and optionally substituted by C₁-C₆-alkyl, C₁-C₆-alkoxy, C₂-C₆-alkenyl, C₂-C₆-alkynyl, amino, acylamino, aminocarbonyl, C₁-C₆-alkoxycarbonyl, aryl, carboxyl, cyano, halogen, hydroxy, nitro, sulfonyl and C₁-C₆-thioalkoxy.

6. (Previously Presented) A sulfonamide compound according to claim 5, wherein Ar¹ is an unsubstituted or substituted phenyl.

7. (Previously Presented) A sulfonamide compound according to claim 5, wherein Ar² is an unsubstituted or substituted thienyl or furanyl group.

8. (Previously Presented) A sulfonamide compound according to claim 1, wherein Ar^1 is selected from the group consisting of a 4-chlorophenyl, nitrophenyl, hydroxyphenyl, alkoxy phenyl, pyridyl, 3,4,-dihydroxyphenyl, thioxo-dihydropyridine or its tautomer and pyrazole and X is O, R^1 is hydrogen, n is 1, Ar^2 is thienyl or furanyl.

9. (Currently Amended) A sulfonamide compound according to claim 8, wherein Y is



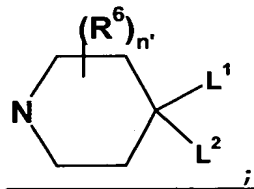
$(\text{R}^6)_n$ is selected from the group consisting of substituted or unsubstituted $\text{C}_1\text{-C}_6$ alkyl, substituted or unsubstituted $\text{C}_1\text{-C}_6$ alkoxy, OH, halogen, nitro, cyano, sulfonyl, and oxo; n is an integer of from 0 to 4; and wherein L^1 and L^2 are independently selected from each other from the group consisting of H, substituted or unsubstituted $\text{C}_1\text{-C}_6$ -aliphatic alkyl, substituted or unsubstituted $\text{C}_2\text{-C}_6$ alkenyl; or L^1 and L^2 are independently selected from the group consisting of substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, aryl- $\text{C}_1\text{-C}_4$ alkyl, heteroaryl- $\text{C}_1\text{-C}_4$ alkyl, $-\text{C}(\text{O})-\text{OR}^3$, $-\text{C}(\text{O})-\text{R}^3$, $-\text{C}(\text{O})-\text{NR}^3\text{R}^3$, $-\text{NR}^3\text{R}^3$; $-\text{NR}^3\text{C}(\text{O})\text{R}^3$, $-\text{NR}^3-\text{C}(\text{O})\text{NR}^3\text{R}^3$, $-(\text{SO})\text{R}^3$, $-(\text{SO}_2)\text{R}^3$, $-\text{NSO}_2\text{R}^3$, $-\text{NHSO}_2\text{R}^3$, and $-\text{SO}_2\text{NR}^3\text{R}^3$;

With R^3 and R^3 being substituents independently selected from the group consisting of H, substituted or unsubstituted $\text{C}_1\text{-C}_4$ alkyl, substituted or unsubstituted $\text{C}_2\text{-C}_6$ alkenyl, substituted or unsubstituted aryl, substituted or

unsubstituted heteroaryl, substituted or unsubstituted aryl-C₁-C₄-alkyl, substituted or unsubstituted heteroaryl-C₁-C₆-alkyl;
said aryl or heteroaryl groups being optionally substituted C₁-C₆-alkyl, C₁-C₆-alkoxy, C₂-C₆-alkenyl, C₂-C₆-alkynyl, amino, acylamino, aminocarbonyl, C₁-C₆-alkoxycarbonyl, aryl, carboxyl, cyano, halogen, hydroxy, nitro, sulfonyl, sulfoxy, C₁-C₆-thioalkoxy.

10. (Currently Amended) A sulfonamide compound according to claim ~~9~~ 8, wherein R⁶ is H, L² is H, L¹ is a 5-membered cyclic group containing 3 heteroatoms; or L¹ is -C(O)-R³, or -NHR³;

wherein Y is



n' is an integer of from 0 to 4;

with R³ being a substituent selected from the group comprising or consisting of C₁-C₄-alkyl, aryl, heteroaryl, aryl-C₁-C₄-alkyl and heteroaryl-C₁-C₆-alkyl;

said aryl or heteroaryl groups being optionally substituted by halogen, hydroxy, nitro, or sulfonyl.

11. (Previously Presented) A sulfonamide compound according to claim 1, selected from the group consisting of:

4-Chloro-N-{5-[4-(3-trifluoromethanesulfonyl-phenylamino)-piperidine-1-sulfonyl]-thiophen-2-ylmethyl}-benzamide;

4-chloro-N-[(5-{[4-(4-fluorobenzoyl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-chloro-N-[(5-{[4-(3-piperidin-1-ylpropyl)piperazin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-chloro-N-({5-[(4-hydroxy-4-phenylpiperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-({5-[(4-benzoylpiperidin-1-yl)sulfonyl]thien-2-yl)methyl}-4-chlorobenzamide;

4-chloro-N-[(5-{[4-(2-oxo-2,3-dihydro-1H-benzimidazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

N-({5-[(4-benzylpiperidin-1-yl)sulfonyl]thien-2-yl)methyl}-4-chlorobenzamide;

4-chloro-N-[(5-{[4-(2,4-difluorobenzoyl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

tert-butyl 1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-ylcarbamate;

4-chloro-N-{[5-(piperidin-1-ylsulfonyl)thien-2-yl)methyl}benzamide;

4-chloro-N-{[5-({3-hydroxy-4-[3-(trifluoromethyl)phenyl]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;

N-[(5-{[4-(benzyloxy)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;

N-(4-chlorophenyl)-2-(5-{[4-(2-oxo-2,3-dihydro-1H-benzimidazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)acetamide;

4-chloro-N-({5-[(4-hydroxypiperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-({5-[(4-benzyl-4-hydroxypiperidin-1-yl)sulfonyl]thien-2-yl)methyl}-4-chlorobenzamide;

N- { [5- ({4- [(2-tert-butyl-1H-indol-5-yl) amino] piperidin-1-yl } sulfonyl) thien-2-yl] methyl } -4-chlorobenzamide;

4-chloro-N- { [5- ({4- [(phenylacetyl) amino] piperidin-1-yl } sulfonyl) thien-2-yl] methyl } benzamide;

N- [(5- { [4- (2H-1,2,3-benzotriazol-2-yl) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] -4-chlorobenzamide;

4-chloro-N- [(5- { [4- (4-chlorobenzoyl) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] benzamide;

4-chloro-N- ({5- [(4-phenoxy piperidin-1-yl) sulfonyl] thien-2-yl } methyl) benzamide;

N- { [5- ({4- [benzyl (methyl) amino] piperidin-1-yl } sulfonyl) thien-2-yl] methyl } -4-chlorobenzamide;

4-chloro-N- { [5- ({4- [3- (2,4-dichlorophenyl) -1H-pyrazol-5-yl] piperidin-1-yl } sulfonyl) thien-2-yl] methyl } benzamide;

4-chloro-N- [(5- { [4- (5-thien-2-yl-1H-pyrazol-3-yl) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] benzamide;

4-chloro-N- [(5- { [4- (2,3,4,5,6-pentamethylbenzoyl) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] benzamide;

4-chloro-N- { [5- ({4- [5- (4-methoxyphenyl) -1H-pyrazol-3-yl] piperidin-1-yl } sulfonyl) thien-2-yl] methyl } benzamide;

N- ({5- [(4-anilinopiperidin-1-yl) sulfonyl] thien-2-yl } methyl) -4-chlorobenzamide;

4-chloro-N- [(5- { [4- (2-phenylethyl) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] benzamide;

N- [(5- { [4- (1H-1,2,3-benzotriazol-1-yl) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] -4-chlorobenzamide;

2- (5- { [4- (1H-1,2,3-benzotriazol-1-yl) piperidin-1-yl] sulfonyl } thien-2-yl) -N- (4-chlorophenyl) acetamide;

2-{1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl}-2H-1,2,3-benzotriazole-5-carboxylic acid;

4-chloro-N-[(5-{[4-(5-chloro-1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

methyl 1-{1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl}-1H-1,2,3-benzotriazole-5-carboxylate;

methyl 1-{1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl}-1H-1,2,3-benzotriazole-6-carboxylate;

methyl 2-{1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl}-2H-1,2,3-benzotriazole-5-carboxylate;

4-chloro-N-[(5-{[4-(6-chloro-1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-chloro-N-{[5-({4-[5-(trifluoromethyl)-1H-1,2,3-benzotriazol-1-yl]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;

N-[(5-{[4-(7-aza-1H-benzimidazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;

1-{1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl}-1H-1,2,3-benzotriazole-5-carboxylic acid;

1-{1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl}-1H-1,2,3-benzotriazole-6-carboxylic acid;

N-[(5-{[4-(2-amino-9H-purin-9-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;

4-chloro-N-[(5-{[4-(9H-purin-9-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

N-[(5-{[4-(6-amino-9H-purin-9-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;
4-chloro-N-[(5-{[4-(6-nitro-1H-benzimidazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(5-nitro-1H-benzimidazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(2H-1,2,3-triazol-2-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
N-[(5-{[4-(1H-benzimidazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;
4-chloro-N-[(5-{[4-[3-propylanilino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-[3-(trifluoromethyl)anilino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-[3-(dimethylamino)anilino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
methyl 3-[(1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl)amino]-benzoate;
4-chloro-N-[(5-{[4-[3-(methylsulfonyl)anilino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-[3-nitroanilino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(2-methoxyanilino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
3-[(1-[(5-{[(4-chlorobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl)amino]benzamide;
4-chloro-N-[(5-{[4-[2-(trifluoromethyl)anilino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-chloro-N-({5-[(4-{2-nitro-4-
[(trifluoromethyl)sulfonyl]anilino}piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;
4-chloro-N-[(5-{[4-(4-chloroanilino)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;
4-chloro-N-{[5-({4-[4-
(trifluoromethyl)anilino]piperidin-1-yl)sulfonyl]thien-2-
yl)methyl}benzamide;
4-chloro-N-({5-[(4-{4-
[(trifluoromethyl)sulfonyl]anilino}piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;
4-chloro-N-({5-[(4-{2-nitroanilino}piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;
N-{[5-({4-[4-(aminocarbonyl)anilino]piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}-4-chlorobenzamide;
4-chloro-N-{[5-({4-[4-(1,3-dithiolan-2-
yl)anilino]piperidin-1-yl)sulfonyl]thien-2-
yl)methyl}benzamide;
N-[(5-{[4-(3-chloroanilino)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl]-3-nitrobenzamide;
4-chloro-N-[(5-{[4-(3-chloroanilino)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;
4-chloro-N-[(5-{[4-(3-methoxyanilino)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;
4-chloro-N-{[5-({4-[3-
(methylsulfonyl)anilino]piperidin-1-yl)sulfonyl]thien-2-
yl)methyl}benzamide;
N-({5-[(4-{3-[amino(imino)methyl]anilino}piperidin-
1-yl)sulfonyl]thien-2-yl)methyl}-4-chlorobenzamide;
4-chloro-N-({5-[(4-{3-[(2-
hydroxyethyl)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-
yl)methyl}benzamide;

N-[(5-{[4-(2-aminoanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;
4-chloro-N-[(5-{[4-(2-hydroxyanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(4-hydroxyanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(3-[(trifluoromethyl)sulfanyl]anilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(3-toluidino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(3-chloro-5-(trifluoromethyl)pyridin-2-yl]amino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(3-(1,3-oxazol-5-yl)anilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
N-[(5-{[4-(3-tert-butylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;
4-chloro-N-[(5-{[4-(2-propylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(2,2-dioxido-1,3-dihydro-2-benzothien-5-yl)amino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(2,3-dihydro-1H-inden-5-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(4-propylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-[(5-{[4-(3-nitropyridin-2-yl)amino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
N-[(5-{[4-(3-aminopyridin-2-yl)amino]piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;

N-[(5-{[4-([1,1'-biphenyl]-3-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;

N-[(5-{[4-(3-benzylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;

4-chloro-N-[(5-{[4-(pyrimidin-2-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-chloro-N-{[5-([4-[4-(morpholin-4-ylsulfonyl)anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

4-chloro-N-([5-([4-([4-(trifluoromethyl)pyrimidin-2-yl]amino)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]benzamide;

4-chloro-N-[(5-{[4-(3-cyclohexyl-4-hydroxyanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

N-([5-([4-3-[(butylamino)sulfonyl]anilino)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-4-chlorobenzamide;

4-chloro-N-[(5-{[4-(3-ethylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-chloro-N-[(5-{[4-(5,6,7,8-tetrahydronaphthalen-1-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

N-{[5-([4-[3-(aminosulfonyl)anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-4-chlorobenzamide;

4-chloro-N-[(5-{[4-(quinolin-5-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-chloro-N-[(5-{[4-(quinolin-8-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-Chloro-N-[(5-{[4-(3-propylphenoxy)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

4-nitro-N-([5-([4-3-[(trifluoromethyl)sulfonyl]anilino)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]benzamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-3-nitrobenzamide;

4-nitro-N-({5-[(4-{3-
[(trifluoromethyl)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-[(5-{[4-(2,4-difluorobenzoyl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-3-nitrobenzamide;

4-nitro-N-({5-[(4-{3-
[(trifluoromethyl)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-[(5-{[4-(2,4-difluorobenzoyl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;

3-nitro-N-[(5-{[4-(3-methoxyanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

3-nitro-N-{[5-({4-[3-
(trifluoromethyl)anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;

N-{[5-({4-[3-(dimethylamino)anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl]-3-nitrobenzamide;

3-nitro-N-{[5-({4-[3-
(methylsulfonyl)anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;

3-nitro-N-{[5-({4-[3-
(methylsulfanyl)anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;

N- { [5- ({4- [3- (aminosulfonyl) anilino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } -3-nitrobenzamide;
methyl 3- { [1- ({5- [({3-nitrobenzoyl} amino) methyl] -thien-2-yl} sulfonyl) -piperidin-4-yl] amino } benzoate;
N- { [5- ({4- [3- (aminocarbonyl) anilino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } -3-nitrobenzamide;
3-nitro-N- ({5- [(4- {3-nitroanilino} piperidin-1-yl) sulfonyl] thien-2-yl} methyl) benzamide;
3-nitro-N- [(5- { [4- (2-methoxyanilino) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] benzamide;
3-nitro-N- { [5- ({4- [2- (trifluoromethyl) anilino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } benzamide;
3-nitro-N- ({5- [(4- {2-nitroanilino} piperidin-1-yl) sulfonyl] thien-2-yl} methyl) benzamide;
N- [(5- { [4- (4-chloroanilino) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] -3-nitrobenzamide;
3-nitro-N- { [5- ({4- [4- (trifluoromethyl) anilino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } benzamide;
3-nitro-N- ({5- [(4- {4- [(trifluoromethyl) sulfonyl] anilino} piperidin-1-yl) sulfonyl] thien-2-yl} methyl) benzamide;
N- { [5- ({4- [4- (aminocarbonyl) anilino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } -3-nitrobenzamide;
N- [(5- { [4- (3-propylanilino) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] -3-nitrobenzamide;
N- [(5- { [4- (3-chloroanilino) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] -4-nitrobenzamide;
4-nitro-N- [(5- { [4- (3-methoxyanilino) piperidin-1-yl] sulfonyl } thien-2-yl) methyl] benzamide;

4-nitro-N-{ [5-({4-[3-(trifluoromethyl)anilino]piperidin-1-yl)sulfonyl}thien-2-yl)methyl}benzamide;

N-{ [5-({4-[3-(dimethylamino)anilino]piperidin-1-yl)sulfonyl}thien-2-yl)methyl}-4-nitrobenzamide;

4-nitro-N-[(5-{[4-(3-propylanilino)piperidin-1-yl)sulfonyl}thien-2-yl)methyl]benzamide;

4-nitro-N-{ [5-({4-[3-(methylsulfonyl)anilino]piperidin-1-yl)sulfonyl}thien-2-yl)methyl}benzamide;

4-nitro-N-{ [5-({4-[3-(methylsulfonyl)anilino]piperidin-1-yl)sulfonyl}thien-2-yl)methyl}benzamide;

N-{ [5-({4-[3-(aminosulfonyl)anilino]piperidin-1-yl)sulfonyl}thien-2-yl)methyl}-4-nitrobenzamide;

methyl 3-{ [1-({5-[(4-nitrobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl}amino}benzoate;

3-{ [1-({5-[(4-nitrobenzoyl)amino]methyl}thien-2-yl)sulfonyl]piperidin-4-yl}amino}benzamide;

4-nitro-N-({5-[(4-{3-nitroanilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

4-nitro-N-[(5-{[4-(2-methoxyanilino)piperidin-1-yl)sulfonyl}thien-2-yl)methyl]benzamide;

4-nitro-N-{ [5-({4-[2-(trifluoromethyl)anilino]piperidin-1-yl)sulfonyl}thien-2-yl)methyl}benzamide;

4-nitro-N-({5-[(4-{2-nitroanilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-[(5-{[4-(4-chloroanilino)piperidin-1-yl)sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;

4-nitro-N-([5-([4-[4-(trifluoromethyl)anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]benzamide;

4-nitro-N-([5-([4-([4-(trifluoromethyl)sulfonyl]anilino)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]benzamide;

N-([5-([4-[4-(aminocarbonyl)anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-4-nitrobenzamide;

N-([5-([4-[4-(1,3-dithiolan-2-yl)anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-4-nitrobenzamide;

N-([5-([4-[3-[amino(imino)methyl]anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-3-nitrobenzamide;

N-([5-([4-[3-[(2-hydroxyethyl)sulfonyl]anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-3-nitrobenzamide;

N-([5-([4-anilinopiperidin-1-yl)sulfonyl]thien-2-yl)methyl]-3-nitrobenzamide;

N-([5-([4-[3-[(2-hydroxyethyl)sulfonyl]anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-4-nitrobenzamide;

N-([5-([4-anilinopiperidin-1-yl)sulfonyl]thien-2-yl)methyl]-4-nitrobenzamide;

N-([5-([4-[3-[amino(imino)methyl]anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-4-nitrobenzamide;

3-nitro-N-([5-([4-[3-([4-(trifluoromethyl)sulfonyl]anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]benzamide;

4-nitro-N-([5-([4-[3-([4-(trifluoromethyl)sulfonyl]anilino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]benzamide;

3-nitro-N-([5-([4-([3-nitropyridin-2-yl)amino]piperidin-1-yl)sulfonyl]thien-2-yl)methyl]benzamide;

N-{[5-({4-[(2,2-dioxido-1,3-dihydro-2-benzothien-5-yl)amino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}-3-nitrobenzamide;

N-[(5-{[4-(2,3-dihydro-1H-inden-5-ylamino)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]-3-nitrobenzamide;

3-nitro-N-[(5-{[4-(2-propylanilino)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]benzamide;

3-nitro-N-[(5-{[4-(4-propylanilino)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]benzamide;

N-[(5-{[4-(3-tert-butylanilino)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]-3-nitrobenzamide;

3-nitro-N-{[5-({4-[3-(1,3-oxazol-5-yl)anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;

3-nitro-N-[(5-{[4-(2-phenylethyl)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]benzamide;

N-({5-[(4-{[3-chloro-5-(trifluoromethyl)pyridin-2-yl]amino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}-3-nitrobenzamide;

N-[(5-{[4-([1,1'-biphenyl]-3-ylamino)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]-3-nitrobenzamide;

N-[(5-{[4-(3-benzylanilino)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]-3-nitrobenzamide;

3-nitro-N-{[5-({4-[3-(morpholin-4-yl)sulfonyl]anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;

3-nitro-N-[(5-{[4-(3-propylphenoxy)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]benzamide;

4-nitro-N-[(5-{[4-(pyrimidin-2-ylamino)piperidin-1-yl}sulfonyl}thien-2-yl)methyl]benzamide;

N-{[5-({4-[(3-aminopyridin-2-yl)amino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}-4-nitrobenzamide;

4-nitro-N-[(5-{[4-({3-nitropyridin-2-yl}amino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
N-[(5-{[4-(2,3-dihydro-1H-inden-5-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;
4-nitro-N-[(5-{[4-(2-propylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-nitro-N-[(5-{[4-(4-propylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
N-[(5-{[4-(3-tert-butylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;
4-nitro-N-{[5-({4-[3-(1,3-oxazol-5-yl)anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;
4-nitro-N-[(5-{[4-(2-phenylethyl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
N-({5-[(4-{[3-chloro-5-(trifluoromethyl)pyridin-2-yl]amino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl)-4-nitrobenzamide;
N-[(5-{[4-([1,1'-biphenyl]-3-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;
N-[(5-{[4-(3-benzylanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-nitrobenzamide;
4-nitro-N-{[5-({4-[3-(morpholin-4-ylsulfonyl)anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;
N-[(5-{[4-(2-aminoanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-3-nitrobenzamide;
3-nitro-N-[(5-{[4-(pyrimidin-2-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
N-{[5-({4-[(3-aminopyridin-2-yl)amino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}-3-nitrobenzamide;

N-({5-[(4-{2-nitro-4-
[(trifluoromethyl)sulfonyl]anilino}piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}-3-methoxybenzamide;
3-nitro-N-({5-[(4-{[4-(trifluoromethyl)pyrimidin-2-
yl]amino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;
N-[(5-{[4-(3-cyclohexyl-4-hydroxyanilino)piperidin-
1-yl)sulfonyl]thien-2-yl)methyl}-3-nitrobenzamide;
N-({5-[(4-{3-
[(butylamino)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-
yl)methyl}-3-nitrobenzamide;
N-[(5-{[4-(3-ethyl-anilino)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}-3-nitrobenzamide;
3-nitro-N-[(5-{[4-(5,6,7,8-tetrahydronaphthalen-1-
ylamino)piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;
4-nitro-N-[(5-{[4-(3-propylphenoxy)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;
N-[(5-{[4-(2,4-difluorobenzoyl)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}-3-nitrobenzamide;
N-[(5-{[4-(2,4-difluorobenzoyl)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}-3-methoxybenzamide;
2-Hydroxy-N-({5-[(4-{3-
[(trifluoromethyl)sulfonyl]anilino}piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;
N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}-3-methoxybenzamide;
N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}-2-hydroxybenzamide;
N-[(5-{[4-{[4-(1,3-dithiolan-2-yl)anilino]piperidin-
1-yl)sulfonyl]thien-2-yl)methyl}-3-nitrobenzamide;
3-methoxy-N-[(5-{[4-(3-methoxyanilino)piperidin-1-
yl)sulfonyl]thien-2-yl)methyl}benzamide;

3-methoxy-N-{ [5- ({4- [3-
(trifluoromethyl) anilino] piperidin-1-yl} sulfonyl) thien-2-
yl] methyl} benzamide;

N- { [5- ({4- [3- (dimethylamino) anilino] piperidin-1-
yl} sulfonyl) thien-2-yl] methyl} -3-methoxybenzamide;

3-methoxy-N- [(5- { [4- (3-propylanilino) piperidin-1-
yl] sulfonyl}) thien-2-yl] methyl] benzamide;

3-methoxy-N- { [5- ({4- [3-
(methylsulfonyl) anilino] piperidin-1-yl} sulfonyl) thien-2-
yl] methyl} benzamide;

3-methoxy-N- { [5- ({4- [3-
(methylsulfonyl) anilino] piperidin-1-yl} sulfonyl) thien-2-
yl] methyl} benzamide;

N- { [5- ({4- [3- (aminosulfonyl) anilino] piperidin-1-
yl} sulfonyl) thien-2-yl] methyl} -3-methoxybenzamide;

methyl 3- ({1- [(5- { [(3-methoxybenzoyl) amino] -
methyl}) thien-2-yl] sulfonyl] piperidin-4-yl} amino) -benzoate;

N- { [5- ({4- [3- (aminocarbonyl) anilino] piperidin-1-
yl] sulfonyl) thien-2-yl] methyl} -3-methoxybenzamide;

3-methoxy-N- [(5- { [4- (2-methoxyanilino) piperidin-1-
yl] sulfonyl}) thien-2-yl] methyl] benzamide;

N- ({5- [(4- {3-nitroanilino} piperidin-1-
yl] sulfonyl]) thien-2-yl] methyl} -3-methoxybenzamide;

3-methoxy-N- { [5- ({4- [2-
(trifluoromethyl) anilino] piperidin-1-yl} sulfonyl) thien-2-
yl] methyl} benzamide;

N- ({5- [(4- {2-nitroanilino} piperidin-1-
yl] sulfonyl]) thien-2-yl] methyl} -3-methoxybenzamide;

N- { [5- ({4- [4- (aminocarbonyl) anilino] piperidin-1-
yl] sulfonyl) thien-2-yl] methyl} -3-methoxybenzamide;

N- { [5- ({4- [4- (1,3-dithiolan-2-yl) anilino] piperidin-
1-yl] sulfonyl) thien-2-yl] methyl} -3-methoxybenzamide;

N-[(5-{[4-(3-chloroanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-3-methoxybenzamide;

N-[(5-{[4-(4-chloroanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-3-methoxybenzamide;

3-methoxy-N-({5-[(4-{4-[(trifluoromethyl)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-({5-[(4-{3-[amino(imino)methyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}-3-methoxybenzamide;

N-({5-[(4-{3-[(2-hydroxyethyl)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}-3-methoxybenzamide;

3-methoxy-N-({5-[(4-{3-[(trifluoromethyl)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-({5-[(4-anilinopiperidin-1-yl)sulfonyl]thien-2-yl)methyl}-3-methoxybenzamide;

3-methoxy-N-({5-[(4-{3-[(trifluoromethyl)sulfanyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-[(5-{[4-(4-hydroxyanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-3-methoxybenzamide;

3-nitro-N-({5-[(4-{3-[(trifluoromethyl)sulfanyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

4-nitro-N-({5-[(4-{3-[(trifluoromethyl)sulfanyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;

N-[(5-{[4-(2-hydroxyanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-3-methoxybenzamide;

3-methoxy-N-[(5-{[4-(pyrimidin-2-ylamino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

N- { [5- ({4- [(3-aminopyridin-2-yl) amino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } -3-methoxybenzamide;

N- [(5- { [4- ({3-nitropyridin-2-yl} amino) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] -3-methoxybenzamide;

N- { [5- ({4- [(2,2-dioxido-1,3-dihydro-2-benzothien-5-yl) amino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } -3-methoxybenzamide;

N- [(5- { [4- (2,3-dihydro-1H-inden-5-ylamino) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] -3-methoxybenzamide;

3-methoxy-N- [(5- { [4- (2-propylanilino) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] benzamide;

3-methoxy-N- [(5- { [4- (4-propylanilino) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] benzamide;

N- [(5- { [4- (3-tert-butylanilino) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] -3-methoxybenzamide;

N- ({5- [(4- { [3-chloro-5- (trifluoromethyl) pyridin-2-yl] amino} piperidin-1-yl) sulfonyl] thien-2-yl} methyl) -3-methoxybenzamide;

3-methoxy-N- { [5- ({4- [3- (1,3-oxazol-5-yl) anilino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } benzamide;

N- [(5- { [4- ([1,1'-biphenyl] -3-ylamino) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] -3-methoxybenzamide;

3-methoxy-N- [(5- { [4- (3-propylphenoxy) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] benzamide;

3-methoxy-N- { [5- ({4- [3- (morpholin-4-ylsulfonyl) anilino] piperidin-1-yl} sulfonyl) thien-2-yl] methyl } benzamide;

3-methoxy-N- [(5- { [4- (2-phenylethyl) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] benzamide;

N- [(5- { [4- (3-benzylanilino) piperidin-1-yl] sulfonyl) thien-2-yl] methyl] -3-methoxybenzamide;

3-methoxy-N-({5-[4-{[4-(trifluoromethyl)pyrimidin-2-yl]amino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl)benzamide;

N-[(5-{[4-(3-cyclohexyl-4-hydroxyanilino)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-3-methoxybenzamide;

N-({5-[4-{3-[(butylamino)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl)-3-methoxybenzamide;

N-[(5-{[4-(3-ethylanilino)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-3-methoxybenzamide;

3-methoxy-N-[(5-{[4-(5,6,7,8-tetrahydronaphthalen-1-ylamino)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]benzamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-5-nitro-1H-pyrazole-3-carboxamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-2-oxo-1,2-dihydropyridine-3-carboxamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-2-thioxo-1,2-dihydropyridine-3-carboxamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-3,4-dihydroxybenzamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]pyridine-2-carboxamide;

N-[(5-{[4-(hexyloxy)piperidin-1-yl)sulfonyl]thien-2-yl)methyl]-3-methoxybenzamide;

N-({5-[4-(heptanoyl)piperidin-1-yl)sulfonyl]thien-2-yl)methyl)-3-methoxybenzamide;

4-chloro-N-[(5-{[4-(3-propylanilino)piperidin-1-yl)sulfonyl]-2-furyl)methyl]benzamide;

4-chloro-N-[(5-{[4-(3-chloroanilino)piperidin-1-yl]sulfonyl}-2-furyl)methyl]benzamide;

4-chloro-N-[(5-{[4-(3-methoxyanilino)piperidin-1-yl]sulfonyl}-2-furyl)methyl]benzamide;

4-chloro-N-{[5-({4-[3-(trifluoromethyl)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}benzamide;

4-chloro-N-{[5-({4-[3-(dimethylamino)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}benzamide;

4-chloro-N-{[5-({4-[3-(methylsulfonyl)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}benzamide;

4-chloro-N-{[5-({4-[3-(methylsulfonyl)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}benzamide;

N-{[5-({4-[3-(aminosulfonyl)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}-4-chlorobenzamide;

methyl 3-({1-[(5-{[(4-chlorobenzoyl)amino]methyl}-2-furyl)sulfonyl]piperidin-4-yl}amino)benzoate;

3-({1-[(5-{[(4-chlorobenzoyl)amino]methyl}-2-furyl)sulfonyl]piperidin-4-yl}amino)benzamide;

4-chloro-N-({5-[(4-{3-nitroanilino}piperidin-1-yl)sulfonyl]-2-furyl)methyl}benzamide;

4-chloro-N-[(5-{[4-(2-methoxyanilino)piperidin-1-yl]sulfonyl}-2-furyl)methyl]benzamide;

4-chloro-N-{[5-({4-[2-(trifluoromethyl)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}benzamide;

4-chloro-N-({5-[(4-{2-nitroanilino}piperidin-1-yl)sulfonyl]-2-furyl)methyl}benzamide;

4-chloro-N-[(5-{[4-(4-chloroanilino)piperidin-1-yl]sulfonyl}-2-furyl)methyl]benzamide;

4-chloro-N-{[5-({4-[4-(trifluoromethyl)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}benzamide;

4-chloro-N-({5-[4-{4-[(trifluoromethyl)sulfonyl]anilino}piperidin-1-yl]sulfonyl}-2-furyl)methyl}benzamide;

N-{[5-({4-[4-(aminocarbonyl)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}-4-chlorobenzamide;

4-chloro-N-{[5-({4-[4-(1,3-dithiolan-2-yl)anilino]piperidin-1-yl}sulfonyl)-2-furyl]methyl}benzamide;

N-({5-[4-{3-[amino(imino)methyl]anilino}piperidin-1-yl]sulfonyl}-2-furyl)methyl)-4-chlorobenzamide;

4-chloro-N-({5-[4-{3-[(trifluoromethyl)sulfonyl]anilino}piperidin-1-yl]sulfonyl}-2-furyl)methyl}benzamide;

N-({5-[4-anilinopiperidin-1-yl]sulfonyl}-2-furyl)methyl)-4-chlorobenzamide; and

4-nitro-N-({5-[4-{3-[(trifluoromethyl)sulfanyl]anilino}piperidin-1-yl]sulfonyl}2-furyl)methyl}benzamide.

12. (Previously Presented) A sulfonamide compound according to claim 11, which is selected from the group consisting of:

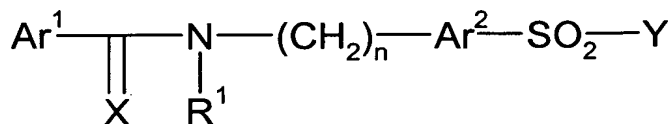
4-chloro-N-[(5-{[4-(2,4-difluorobenzoyl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;

N-({5-[4-anilinopiperidin-1-yl]sulfonyl}thien-2-yl)methyl)-4-chlorobenzamide;

N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;

N-[(5-{[4-(1H-benzimidazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-4-chlorobenzamide;
4-chloro-N-{[5-({4-[3-propylanilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl}benzamide;
4-chloro-N-[(5-{[4-(4-chloroanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
4-chloro-N-({5-[(4-{3-[(2-hydroxyethyl)sulfonyl]anilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}benzamide;
N-{[5-({4-[3-(aminosulfonyl)anilino]piperidin-1-yl}sulfonyl)thien-2-yl)methyl]-4-chlorobenzamide;
4-nitro-N-[(5-{[4-(3-methoxyanilino)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]benzamide;
methyl 3-{[1-({5-[(4-nitrobenzoyl)amino)methyl]thien-2-yl}sulfonyl)piperidin-4-yl]amino}benzoate;
N-[(5-{[4-(1H-1,2,3-benzotriazol-1-yl)piperidin-1-yl]sulfonyl}thien-2-yl)methyl]-2-hydroxybenzamide; and
N-({5-[(4-{2-nitroanilino}piperidin-1-yl)sulfonyl]thien-2-yl)methyl}-3-methoxybenzamide.

13. (Currently Amended) A method for the modulation of the JNK pathway, comprising administering an effective amount of a sulfonamide compound according to formula I



I

wherein Ar¹ and Ar² are independently from each other substituted or ~~unsubstituted~~ unsubstituted aryl or heteroaryl groups;

X is O or S;

R¹ is hydrogen or a C₁-C₆-alkyl group, or R¹ forms a substituted or unsubstituted 5-6-membered saturated or unsaturated ring with Ar¹;

n is an integer from 0 to 5;

Y within formula I is a piperidino group; whereby one nitrogen within said piperidino group forms a bond with the sulfonyl group of formula I, thus providing a sulfonamide.

14. (Previously Presented) A method according to claim 13 for the treatment of disorders associated with the abnormal expression or activity of JNK.

15. (Previously Presented) A method according to claim 14 for the treatment of disorders associated with abnormal expression or activity of JNK2 and/or 3.

16. (Previously Presented) A method for the treatment of neuronal disorders comprising administering to a patient in need thereof an effective amount of a sulfonamide compound according to formula I in claim 1.

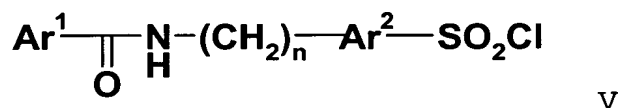
17. (Currently Amended) A method for the treatment of autoimmune diseases comprising administering to a patient in need thereof an effective amount of a sulfonamide compound according to formula I in claim ~~13~~1.

18. (Currently Amended) A method for the treatment of cancer comprising administering to a patient in need thereof an effective amount of a sulfonamide compound according to formula I in claim ~~13~~1.

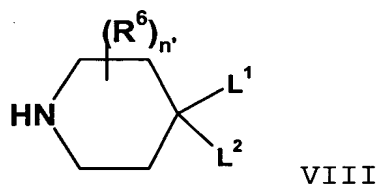
19. (Currently Amended) A method for the treatment of cardiovascular diseases comprising administering to a patient in need thereof an effective amount of a sulfonamide compound according to formula I in claim ~~13~~1.

20. (Previously Presented) A pharmaceutical composition containing at least one sulfonamide compound according to claim 1 and a pharmaceutically acceptable carrier, diluent or excipient.

21. (Previously Presented) Process for the preparation of a sulfonamide compound according to claim 1, comprising reacting a sulfonyl chloride V



with an amine VIII

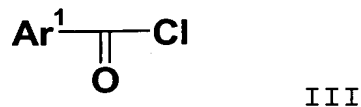


whereby $(\text{R}^6)_n$, L^1 and L^2 are as defined in claim 1.

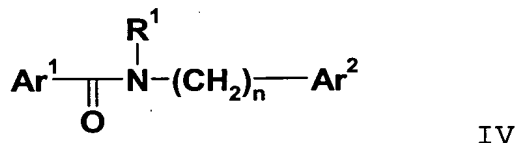
22. (Currently Amended) A process according to claim 21, wherein said sulfonyl chloride V is obtained by
a) coupling an amine of formula II:



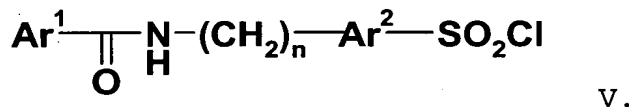
with an acyl chloride of formula III:



where ~~Ar¹ is as defined in claim 1,~~ to provide an amide of formula IV:



b) sulfonating the amide of formula IV to provide a sulfonyl chloride V



23. (Previously Presented) A method in accordance with claim 16, wherein said neuronal disorder is selected from the group consisting of epilepsy, Alzheimer's disease, Huntington's disease, Parkinson's disease, retinal diseases, spinal cord injury, and head trauma.

24. (Previously Presented) A method in accordance with claim 17, wherein said autoimmune disease is selected from the group consisting of multiple sclerosis, inflammatory bowel disease (IBD), rheumatoid arthritis, asthma, septic shock, and transplant rejection.

25. (Previously Presented) A method in accordance with claim 18, wherein said cancer is selected from the group consisting of breast-, colorectal-, and pancreatic cancer.

26. (Previously Presented) A method in accordance with claim 19, wherein said cardiovascular disease is selected from the group consisting of stroke, arteriosclerosis, myocardial infarction, and myocardial reperfusion injury.

27. (Previously Presented) Sulfonamide compounds according to claim 1 wherein n is an integer of from 1 to 3.

28. (Previously Presented) Sulfonamide compounds according to claim 27 wherein n is 1.

29. (Previously Presented) Compositions according to claim 2 wherein n is an integer of from 1 to 3.

30. (Previously Presented) The composition according to claim 29 wherein n is 1.

31. (Previously Presented) Compositions according to claim 3 wherein n is 1 or 2.

32. (Previously Presented) The method according to claim 13 wherein X is O.

33. (Previously Presented) The method according to claim 13 wherein n is an integer of from 1 to 3.

34. (Previously Presented) The method according to claim 33 wherein n is 1.

35. (Previously Presented) The sulfonamide compound according to claim 5 wherein the C₁-C₄ alkyl group is trihalomethyl.

36. (Previously Presented) The sulfonamide compound according to claim 10 wherein L¹ is a triazole ring

which is fused with an unsubstituted or substituted aryl or heteroaryl.

37. (Previously Presented) A method for treating diseases or disorders which are mediated by the JNK function or pathways comprising administering to a patient in need thereof a compound according to claim 1 that inhibits the JNK function of pathways.

38. (Previously Presented) The method according to claim 37 wherein the disease or disorder is a neuronal disorder selected from the group consisting of epilepsy, Alzheimer's disease, Huntington's disease, Parkinson's disease, retinal diseases, spinal cord injury, and head trauma.

39. (Previously Presented) The method according to claim 37 wherein the disease is an autoimmune disease selected from the group consisting of multiple sclerosis, inflammatory bowel disease (IBD), rheumatoid arthritis, asthma, septic shock, and transplant rejection.

40. (Previously Presented) The method according to claim 37 wherein the disease is cancer selected from the group consisting of breast, colorectal, and pancreatic cancer.

41. (Previously Presented) The method according to claim 37 wherein the disease is a cardiovascular disease selected from the group consisting of stroke, arteriosclerosis, myocardial infarction, and myocardial reperfusion injury.